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| (54) Title: ALKOXYI ATE OF 2-PROPYL HEPTA | | | · |

(54) Title: ALKOXYLATE OF 2-PROPYL HEPTANOL AND USE THEREOF

(57) Abstract

The invention relates to an alkoxylate of the general formula (I): $C_5H_{11}CH(C_3H_7)CH_2O(B)_r(C_2H_4O)_pH$, wherein B is an alkyleneoxy group having 3-4 carbon atoms, p is 1-10 and r is 1-6. The alkoxylate may be included as a surface-active component in compositions for cleaning textile materials.

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ALKOXYLATE OF 2-PROPYL HEPTANOL AND USE THEREOF

The present invention relates to an alkoxylate of 2-propyl heptanol. The alkoxylate exhibits high detergent 5 power on textile materials and low foaming compared with similar compounds having a hydrophobic group of approximately the same size and approximately the same HLB-value. The alkoxylate may advantageously be used as a surfaceactive component in detergent compositions for textile materials.

It has long been known to alkoxylate alcohols for obtaining non-ionic surface-active compounds. These compounds have been used in detergent compositions because of their wetting and dispersing properties. In a number of applications, alkoxylates of C_{8-11} alcohols have how-15 ever been found to be too high-foaming and/or not to have the desired detergent power. For example, ethoxylates based on branched C_8 alcohols often exhibit acceptable foaming but too low a detergent power, whereas ethoxylates based on straight or branched alcohols having 20 a larger hydrocarbon chain often show an acceptable surface activity but too high foaming. Thus, there is a need for new alkylene oxide adducts with an improved ratio of foaming to detergent power.

It has now been found that an alkoxylate based on 2-propyl heptanol has good detergent and wetting properties as well as low foaming as compared with other alcohols having substantially the same chain length. In addition, it has been found that the alkoxylate is easily degradable and has a surprisingly low biotoxicity. In tests, no skin-irritant effect has been noted.

The alkoxylate according to the invention can be illustrated by the formula

$$^{35} \quad ^{C_5H_{11}CH(C_3H_7)CH_2O(B)_r(C_2H_4O)_pH}$$
 (I)

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wherein B is an alkyleneoxy group having 3-4 carbon atoms, p is 1-10 and r is 1-6. Preferably, p is 2-8 and r is 1-4. In these compounds, the hydrophobic properties of the hydrocarbon chain have been enhanced by adding hydrophobic 5 alkyleneoxy groups closest to the alcohol. The compounds have a good detergent power on textile materials while at the same time showing slightly lower foaming in relation to compounds having a hydrophobic group of approximately the same hydrophobicity and approximately the same HLB-value.

The alkoxylates according to the invention described above can be prepared by adding in a conventional manner in the presence of a conventional alkali catalyst, such as potassium hydroxide or sodium hydroxide, the above-men-15 tioned amounts of alkylene oxide to 2-propyl heptanol, which is a so-called Guebert alcohol. According to a preferred mode of execution, the addition of ethylene oxide is performed using a conventional catalyst which gives a narrower distribution of added ethylene oxide than 20 any alkali catalyst, such as NaOH or KOH. Thus prepared alkoxylates according to the invention have very low foaming. Examples of conventional catalysts giving a narrow distribution of added alkylene oxide are Ca(OH)2, Ba(OH)2, Sr(OH), and hydrotalcite. The reaction is preferably conducted in the absence of free water to reduce the amount of by-products and usually at a temperature of 70-180°C.

Textile-cleaning compositions including the alkoxylate according to the invention may also contain other surface-active compounds, such as anionic ones. Examples 30 hereof are alkyl sulphate, alkyl ether sulphate, alkyl benzene sulphonate, α -olefin sulphonate and alkyl glyceryl sulphonate. Other commonly occurring components are solutising additives, complexing agents and/or pH-adjusting agents, enzymes, bactericides and perfumes. The compositions are usually aqueous and in the form of emulsions, microemulsions or solutions.

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The invention will be further illustrated by the following Examples.

Example 1

An alkoxylate according to the invention are prepared by alkoxylating 2-propyl ethanol with the amounts of alkylene oxide appearing from the Table below in the presence of potassium hydroxide as catalyst. For reference purposes, two alkoxylates were prepared using a C_{9-11} alcohol (Dobanol 91 Shell) as hydrophobic ingredient. The resulting products were analysed and structurally determined by gas chromatography and mass spectrometry. The turbidity points were measured in water or monobutylether diethylene glycol. The following results were obtained.

Table 1

| Com- pound | Alcohol | Mole of alkylene oxide/mole of alcohol | Cata- lyst | Turbidi point | ty |
|---------------|---------------------------|--|---------------|------------------|-----|
| | | or arconor | | Water | BDG |
| 1 | 2-propyl heptanol | 4 PO+6 EO ¹⁾ | кон | 25 | _ |
| A | C ₉₋₁₁ alcohol | 4 EO | кон | - | 62 |
| В | C ₉₋₁₁ alcohol | 6 EO | кон | 56 | _ |

EO = ethylene oxide; PO = propylene oxide, BDG = monobutylether diethylene glycol

1) PO added first

Example 2

The foaming properties of the alkoxylates reported in the following Table were measured according to Ross-Miles ASTM D 1173-53. The following results were obtained.

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Table 2

| Compound | Foam he: | ight, cm |
|----------|----------|----------|
| | O min | 5 min |
| 1 | 83 80 | 12 20 |
| . в | 95 | 30 |

The compound according to the invention has equivalent or slightly lower foaming as compared with compounds

Example 3

A and B.

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Washing tests were carried out in a Terg-O-Tometer on pigment-soiled cotton and cotton/polyester. Washed-away soil was thereafter determined by conventional reflectance measurement. The following results were obtained.

Table 3

| | Washed-away pigment soil , % | | | | |
|----------|------------------------------|------|-----------|--|--|
| Compound | Cotton Cotton/ | | oolyester | | |
| | 40°C | 40°C | 60°C | | |
| 1 | 78 | 73 | 66 | | |
| A | 78 | 65 | 52 | | |

From these results it appears that the compound according to the invention all in all has higher detergent power than the reference compound. From Example 2 also appears that the compound according to the invention has slightly lower foaming than the reference compound.

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CLAIMS

An alkoxylate as claimed in claim 1 or 2, c h a r a c t e r i s e d by having the general formula

$$C_5H_{11}CH(C_3H_7)CH_2O(B)_r(C_2H_4O)_pH$$
 (I)

wherein B is an alkyleneoxy group having 3-4 carbon atoms, 10 p is 1-10 and r is 1-6.

- 2. An alkoxylate as claimed in claim 1, characterised in that p is 2-8 and r is 1-4.
- 15 3. The use of a compound as claimed in claim 1 or 2 in a detergent composition for textile materials.

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CLASSIFICATION OF SUBJECT MATTER IPC5: C07C 43/11, C11D 1/72 According to International Patent Classification (IPC) or to both national classification and IPC **B. FIELDS SEARCHED** Minimum documentation searched (classification system followed by classification symbols) IPC5: C07C, C11D Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched SE,DK,FI,NO classes as above Electronic data base consulted during the international search (name of data base and, where practicable, search terms used) C. DOCUMENTS CONSIDERED TO BE RELEVANT Relevant to claim No. Citation of document, with indication, where appropriate, of the relevant passages Category* 1-3 GB, A, 2145726 (DIVERSEY CORPORATION), Α 3 April 1985 (03.04.85) 1-3 CH, A5, 674358 (OUHADI TRAZOLLAH ET AL.), 31 May A 1990 (31.05.90) 1-3 EP, A1, 0046582 (CONOCO INC.), 3 March 1982 Α (03.03.82)1-3 US, A, 3567784 (WILLIAM T. TSATSOS ET AL.), 2 March 1971 (02.03.71) See patent family annex. Further documents are listed in the continuation of Box C. l x later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention Special categories of cited documents: "A" document defining the general state of the art which is not considered to be of particular relevance "X" document of particular relevance: the claimed invention cannot be "E" ertier document but published on or after the international filing date considered novel or cannot be considered to involve an inventive document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified) step when the document is taken alone document of particular relevance: the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination document referring to an oral disclosure, use, exhibition or other being obvious to a person skilled in the art document published prior to the international filing date but later than "&" document member of the same patent family the priority date claimed Date of mailing of the international search report Date of the actual completion of the international search 22 February 1994 Authorized officer Name and mailing address of the ISA/ Swedish Patent Office Box 5055, S-102 42 STOCKHOLM Eva Johansson Facsimile No. +46 8 666 02 86 Telephone No. +46 8 782 25 00



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A(10-E8A, 12-W12C) B(4-C3C, 10-E4C, 12-M9) C(4-F) F(3-C5, 5-A6D) G(2-A3) N(4-A, 7-B, 7-D) .3 NOVELTY BADI 2002.04.26 *WO 2003091191-A1 2002.09.18 2002-1043360(+2002DE-1018754)\(2003.11.06) C07C A25 B07 C07 D25 E17 F06 G02 (A97 D18 D21 F09 G03) 43/11, 41/03, C11D 1/72 2003-903407/82 BASF AG

cosmetic, New and known 2-propyl-heptan-1-ol alkoxylates, used e.g. in cleaning, coating, adhesive, textile treatment, cosmetic, pharmaceutical or plant protection formulations, contain ethylene oxide and other alkylene oxide groups (Ger)

MN MW MX MZ NI NO NZ OM PH PL PT RO RU SC SD SE SG SK SL TJ TM TN TR TT TZ UA UG US UZ KP KR KZ LC LK LR LS LT LU LV MA MD MG MK VC VN YU ZA ZM ZW) R(AT BE BG CH CY CZ DE DK EA EE ES FI FR GB GH GM GR HU IE IT KE LS CA CH CN CO CR CU CZ DE DK DM DZ EC EE ES FI GB GD GE GH GM HR HU ID IL IN IS JP KE KG **C2003-256873** N(AE AG AL AM AT AU AZ BA BB BG BR BY BZ LU MC MW MZ NL OA PT RO SD SE SI SK SL SZ

2003.04.25 2003WO-EP04334, 2002.09.18 2002DE-1043360 RULAND A, SCHOLTISSEK M, OETTER G, TAEGER K FR TZ UG ZM ZW) Addnl. Data:

C3C, 10-E4C, 12-M9) D(7-B, 8-B13, 11-A3A) E(10-H1D, 11-D, 11-

2-Propyl-heptan-1-ol alkoxylates (IA) with 3-14 ethylenoxy and 1.2-1.8 propylenoxy or pentenoxy or 1-1.8 butenoxy groups in the molecule are new.

DETAILED DESCRIPTION

2-Propyl-heptan-1-ol alkoxylates (IA) of formula (I), with specified numbers of alkoxylate groups, are new. $C - O - (A)_n - (CH_2CH_2O)_m - H$

A = propylenoxy, butenoxy or pentenoxy; n = 1.2-1.8 or, if A is butenoxy, 1-1.8; and

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1 = 3-14

INDEPENDENT CLAIMS are also included for the following:
(1) use of a wider range of alkoxylates of formula (I; where n = 1-8; m = 2-20) as emulsifiers, foam regulators and wetting agents for hard

- (2) preparation of alkoxylates (IA) by reacting the alkanol with propylene oxide and then with ethylene oxide under alkoxylation conditions, which can be carried out in the presence of a double metal cyanide compound as catalyst;
 - (3) preparation of the alkanol by alkaline dimerization of valeraldehyde to an α,β-unsaturated aldehyde, followed by hydrogenation; and
- (4) cleaning, wetting, coating, adhesive, leather degreasing, humectant, textile treatment or cosmetic, pharmaceutical or plant protection formulations containing (I).

ISF

Alkoxylates (I) are used in surfactant formulations for cleaning hard surfaces, in humectants, cosmetic, pharmaceutical and plant protection formulations, lacquers, coating compositions, adhesives, leather degreasing agents, water treatment, the paper industry, fermentation and mineral processing and in emulsion polymerization

(all claimed). Their uses include humectants for the printing industry; lacquers, coating compositions, paints, pigment formulations and adhesives in the lacquer and film industries; leveling agent or yarn cleaning formulations for the textile industry; fiber processing and treatment for the paper and pulp industry; metal processing, e.g. finishing and electroplating; the food industry; water treatment and drinking water recovery; fermentation; mineral processing and dust control; building ancillaries; preparation of dispersions; and coolants and lubricants.

<u>ADVANTAGE</u>

Alkoxylates (I) have good emulsifying, foam regulation and wetting properties, with low angle of contact on hard surfaces, and reduce the surface tension of liquid systems. (I) have a favorable ecological profile, i.e. are not aquatoxic (EC 50 value over 10 mg/l for algae, daphnia or fish); are readily degradable; and are more compatible with the environment and skin than usual.

EXAMPLE

A mixture of 790 g 2-propylheptanol (2-PH) and 8.5 g potassium hydroxide (45% in water) in an autoclave was dried at 80°C and about

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distilled water) was (A) 10, (B) 15, (C) 17 seconds; and the residual 2comprising 2-PH + 8 EO, was prepared by reacting 2-PH directly with 8 mole EO at 145-155°C; and control (C), comprising 2-PH + 8 EO + 1.5 PO, by reacting 2-PH first with 8 mole EO at 145-155°C and then the values for (D) demineralized water, were (A) 48/32, (B) 48/46, (C) 96/95° on polyethylene; and (A) 37/20, (B) 38/25, (C) 39/24, (D) 41/39° on glass. For wetting cotton (to EN 1/52; 0.2 g/l, 2 g/l soda in reacted completely at elevated pressure, then 1470 ml ethylene oxide seconds at a concentration of 0.2 g/l in water at 40°C, compared with 46/40, (D) 65/64° on V2A steel; (A) 57/40, (B) 65/64, (C) 58/54, (D) (EO) were introduced at 145-155°C and reacted at elevated pressure. 20 mbar. To prepare alkoxylate (A), comprising 2-PH + 1.5 PO + 6 catalyst was neutralized with 3.8 ml glacial acetic acid. Control (B), mN/m with hexadecane and (A) 5.2, (B) 8.4, (C) 8.0 with olive oil. with 1.5 mole PO at 120-130°C. The angles of contact after 0.1/10 After purging with inert gas and cooling to room temperature, the PH content was (A) 0.78, (B) 4.3, (C) 1.9 g/100 g. The interfacial tension (1 g/l, 25°C, 10 minutes) was (A) 7.7, (B) 13.2, (C) 13.9 EO, 518 ml propylene oxide (PO) were added at 120-130°C and

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DEFINITIONS

Preferred Definitions:
n = 1.3-1.7 in (IA) and 1-6 in (I);
m = 3-12 in (IA) and 3-14 in (I); and
C₅H₁₁ = n-C₅H₁₁, C₂H₅CH(CH₃)CH₂ and/or CH₃CH(CH₃)CH₂CH₂.
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